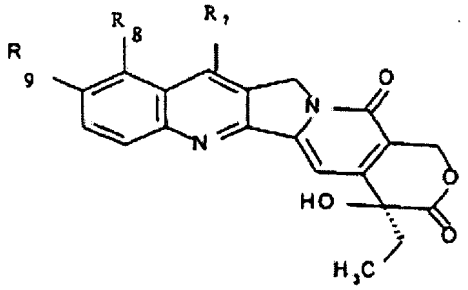
AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the application:

1.-70. (Canceled).

71. (Previously Presented) A method of intracellular delivery of taxol or a camptothecin derivative of formula



wherein: R_7 is a $-C(R_{11})=N-O_{(n)}R_{10}$ group, wherein R_{10} is hydrogen or a C_1-C_5 alkyl or C_2-C_5 alkenyl group, linear or branched or C_3-C_{10} cycloalkyl, group or a linear or branched (C_3-C_{10}) cycloalkyl - (C_1-C_5) alkyl group, or C_6-C_{14} aryl, or a linear or branched (C_6-C_{14}) aryl - (C_1-C_5) alkyl group, or a heterocyclic or linear or branched heterocyclo - (C_1-C_5) alkyl group, said heterocyclic group containing at least a heteroatom selected from the group consisting of nitrogen atom, optionally substituted with a (C_1-C_5) alkyl group, and/or oxygen and/or sulfur; said alkyl, alkenyl, cycloalkyl, cycloalkylalkyl, aryl, aryl-alkyl, heterocyclic or heterocyclo-alkyl groups, being optionally substituted with other groups selected from the group consisting of: halogen, hydroxy, C_1-C_5 alkyl, C_1-C_5 alkoxy, phenyl, cyano, nitro, $-NR_{12}R_{13}$, wherein R_{12} and R_{13}, which may be the same or different, are hydrogen, linear or branched (C_1-C_5) alkyl; a

pharmaceutically acceptable ester of the –COOH group; or the–$CONR_{14}R_{15}$ group, wherein R_{14} and R_{15}, which may be the same or different, are hydrogen or linear or branched (C_1-C_5) alkyl; or

R_{10} is a (C_6-C_{10}) aroyl residue optionally substituted by one or more groups selected from the group consisting of: halogen, hydroxy, linear or branched (C_1-C_5) alkyl, C_1-C_5 alkoxy, phenyl, cyano, nitro, -$NR_{16}R_{17}$, wherein R_{16} and R_{17}, which may be the same or different, is hydrogen, linear or branched (C_1-C_8) alkyl;

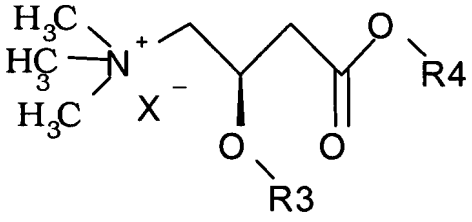
N is the number 0 or 1;

R_{11} is hydrogen, linear or branched C_1-C_5 alkyl, linear or branched C_2-C_5 alkenyl, C_3-C_{10} cycloalkyl, (C_3-C_{10}) cycloalkyl - linear or branched (C_1-C_5) alkyl, C_6-C_{14} aryl, (C_6-C_{14}) aryl - linear or branched alkyl (C_1-C_5);

R_8 and R_9, which may be the same or different are hydrogen, hydroxy, linear or branched C_1-C_5 alkoxy;

their N_1-oxides, their single isomers, in particular the syn and anti isomers of the–$C(R_{11})$=N-$O_{(n)}R_{10}$ group, their possible enantiomers, diastereoisomers and relative admixtures, the pharmaceutically acceptable salts thereof;

using a liposome comprising a compound of formula (II)



(II)

where:

R_3 is a saturated linear or branched acyl chain, with 4-26 carbon atoms;

R_4 is a saturated or unsaturated, linear or branched alkyl chain, with 4-26 carbon atoms; and

X^- is the anion of a pharmacologically acceptable acid.

72. (Previously Presented) The method according to claim 71, in which R_3 is selected from the group consisting of nonanoyl, dodecanoyl, myristoyl, palmitoyl, stearoyl and oleolyl.

73. (Previously Presented) The method according to claim 71, in which R_4 is selected from the group consisting of nonyl, undecyl, tetradecyl, hexadecyl and oleyl.

74. (Previously Presented) The method according to claim 71, in which X^- is selected from the group consisting of chloride; bromide; iodide; aspartate; acid aspartate; citrate; acid citrate; tartrate; acid tartrate; phosphate; acid phosphate; fumarate; acid fumarate; glycerophosphate; glucose phosphate; lactate; maleate; acid maleate; mucate; orotate; oxalate; acid oxalate; sulphate; acid sulphate; trichloroacetate; trifluoroacetate; methane sulphonate; pamoate and acid pamoate.

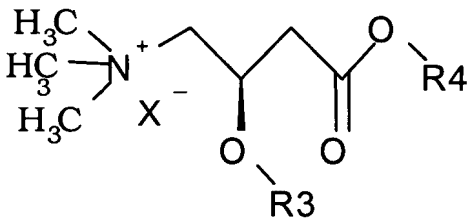
75. (Previously Presented) The method according to claim 71, in which the camptothecin is selected from the group consisting of palmitoyl L-carnitine chloride undecyl ester; stearoyl L-carnitine chloride undecyl ester; stearoyl L-carnitine chloride tetradecyl ester; palmitoyl L-carnitine chloride tetradecyl ester; myristoyl L-carnitine chloride tetradecyl ester; palmitoyl L-carnitine bromide hexadecyl ester, and oleolyl L-carnitine chloride oleyl ester.

76. (Previously Presented) The method according to claim 71, in which said derivative of camptothecin is selected from the group consisting of 7-benzyloxyiminomethylcamptothecin and 7-butoxyiminomethylcam-ptothecin.

77. (Previously Presented) The method according to claim 71, in which the liposome additionally contains helper lipids.

78. (Previously Presented) The method according to claim 77, in which said helper lipid is selected from the group consisting of cholesterol, 1-palmitoyl-2-oleoyl phosphatidyl choline and dioleyl phosphatidyl choline.

79. (Previously Presented) A method of intracellular delivery of a cosmetic using a liposome comprising a compound of formula (II)



(II)

where:

R_3 is a saturated linear or branched acyl chain, with 4-26 carbon atoms;

R_4 is a saturated or unsaturated, linear or branched alkyl chain, with 4-26 carbon atoms; and

X^- is the anion of a pharmacologically acceptable acid.

80. (Previously Presented) The method according to claim 79, in which R_3 is selected from the group consisting of nonanoyl, dodecanoyl, myristoyl, palmitoyl, stearoyl and oleolyl.

81. (Previously Presented) The method according to claim 79, in which R_4 is selected from the group consisting of nonyl, undecyl, tetradecyl, hexadecyl and oleyl.

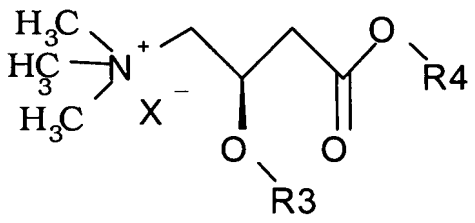
82. (Previously Presented) The method according to claim 79, in which X^- is selected from the group consisting of chloride; bromide; iodide; aspartate; acid aspartate; citrate; acid citrate; tartrate; acid tartrate; phosphate; acid phosphate; fumarate; acid fumarate; glycerophosphate; glucose phosphate; lactate; maleate; acid maleate; mucate; orotate; oxalate; acid oxalate; sulphate; acid sulphate; trichloroacetate; trifluoroacetate; methane sulphonate; pamoate and acid pamoate.

83. (Previously Presented) The method according to claim 79, in which the compound is selected from the group consisting of: palmitoyl L-carnitine chloride undecyl ester; stearoyl L-carnitine chloride undecyl ester; stearoyl L-carnitine chloride tetradecyl ester; palmitoyl L-carnitine chloride tetradecyl ester; myristoyl L-carnitine chloride tetradecyl ester; palmitoyl L-carnitine bromide hexadecyl ester, and oleolyl L-carnitine chloride oleyl ester.

84. (Previously Presented) The method according to claim 79, in which the liposome additionally contains helper lipids.

85. (Previously Presented) The method according to claim 84, in which said helper lipid is selected from the group consisting of cholesterol, 1-palmitoyl-2-oleolyl phosphatidyl choline and dioleoly phospatidyl choline.

86. (Previously Presented) A composition comprising a liposome comprising a compound of formula (II)



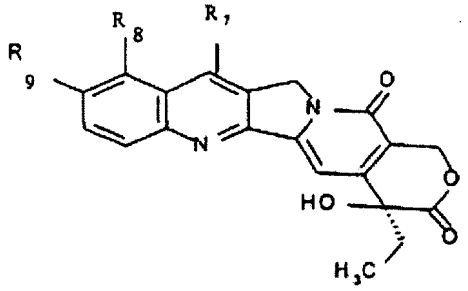
(II)

where:

R_3 is a saturated linear or branched acyl chain, with 4-26 carbon atoms;

R_4 is a saturated or unsaturated, linear or branched alkyl chain, with 4-26 carbon atoms; and

X^- is the anion of a pharmacologically acceptable acid, said liposome comprising taxol or a camptothecin derivative of formula



wherein: R_7 is a $-C(R_{11})=N-O_{(n)}R_{10}$ group, wherein R_{10} is hydrogen or a C_1-C_5 alkyl or C_2-C_5 alkenyl group, linear or branched or C_3-C_{10} cycloalkyl, group or a linear or branched (C_3-C_{10}) cycloalkyl - (C_1-C_5) alkyl group, or C_6-C_{14} aryl, or a linear or branched (C_6-C_{14}) aryl - (C_1-C_5) alkyl group, or a heterocyclic or linear or branched heterocyclo - (C_1-C_5) alkyl group, said heterocyclic group containing at least a heteroatom selected from the group consisting of nitrogen atom, optionally substituted with a (C_1-C_5) alkyl group, and/or oxygen and/or sulfur; said alkyl, alkenyl, cycloalkyl, cycloalkylalkyl, aryl, aryl-alkyl, heterocyclic or heterocyclo-alkyl groups, being optionally substituted with other groups selected from the group consisting of: halogen, hydroxy, C_1-C_5 alkyl, C_1-C_5 alkoxy, phenyl, cyano, nitro, $-NR_{12}R_{13}$, wherein R_{12} and R_{13}, which may be the same or different, are hydrogen, linear or branched (C_1-C_5) alkyl; a pharmaceutically acceptable ester of the –COOH group; or the–$CONR_{14}R_{15}$ group, wherein R_{14} and R_{15}, which may be the same or different, are hydrogen or linear or branched (C_1-C_5) alkyl; or

R_{10} is a (C_6-C_{10}) aroyl residue optionally substituted by one or more groups selected from the group consisting of: halogen, hydroxy, linear or branched (C_1-C_5) alkyl, C_1-C_5 alkoxy, phenyl, cyano, nitro, -$NR_{16}R_{17}$, wherein R_{16} and R_{17}, which may be the same or different, are hydrogen, linear or branched (C_1-C_8) alkyl;

n is the number 0 or 1;

R_{11} is hydrogen, linear or branched C_1-C_5 alkyl, linear or branched C_2-C_5 alkenyl, C_3-C_{10} cycloalkyl, (C_3-C_{10}) cycloalkyl - linear or branched (C_1-C_5) alkyl, C_6-C_{14} aryl, (C_6-C_{14}) aryl - linear or branched alkyl (C_1-C_5);

R_8 and R_9, which may be the same or different is hydrogen, hydroxy, linear or branched C_1-C_5 alkoxy;

their N_1-oxides, their single isomers, in particular the syn and anti isomers of the– $C(R_{11})$=N-$O_{(n)}R_{10}$ group, their possible enantiomers, diastereoisomers and relative admixtures, the pharmaceutically acceptable salts thereof; or

said liposome comprising a substance with cosmetic activity.

87. (Previously Presented) The composition according to claim 86, in which R_3 is selected from the group consisting of nonanoyl, dodecanoyl, myristoyl, palmitoyl, stearoyl and oleolyl.

88. (Previously Presented) The composition according to claim 86, in which R_4 is selected from the group consisting of nonyl, undecyl, tetradecyl, hexadecyl and oleyl.

89. (Previously Presented) The composition according to claim 86, in which X– is selected from the group consisting of chloride; bromide; iodide; aspartate; acid aspartate; citrate; acid citrate; tartrate; acid tartrate; phosphate; acid phosphate; fumarate; acid fumarate; glycerophosphate; glucose phosphate; lactate; maleate; acid maleate; mucate; orotate; oxalate;

acid oxalate; sulphate; acid sulphate; trichloroacetate; trifluoroacetate; methane sulphonate; pamoate and acid pamoate.

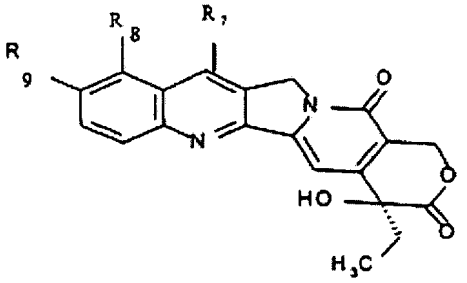
90. (Previously Presented) The composition according to claim 86, in which the compound is selected from the group consisting of: palmitoyl L-carnitine chloride undecyl ester; stearoyl L-carnitine chloride undecyl ester; stearoyl L-carnitine chloride tetradecyl ester; palmitoyl L-carnitine chloride tetradecyl ester; myristoyl L-carnitine chloride tetradecyl ester; palmitoyl L-carnitine bromide hexadecyl ester, and oleolyl L-carnitine chloride oleyl ester.

91. (Previously Presented) The composition according to according to claim 86, in which the liposome additionally contains helper lipids.

92. (Previously Presented) The composition according to claim 91, in which said helper lipid is selected from the group consisting of cholesterol, 1-palmitoyl-2-oleoyl phosphatidyl choline or dioleyl phosphatidyl choline.

93. (Previously Presented) The composition according to claim 86, which composition is administered orally, parenterally, intravenously, intramuscularly, subcutaneously, transdermally or in the form of a nasal or mouth spray.

94. (Previously Presented) A method of transporting an antitumor drug to the target organ of a subject in need of antitumor treatment, wherein said drug is selected from the group consisting of taxol or a camptothecin derivative of formula



wherein: R_7 is a $-C(R_{11})=N-O_{(n)}R_{10}$ group, wherein R_{10} is hydrogen or a C_1-C_5 alkyl or C_2-C_5 alkenyl group, linear or branched or C_3-C_{10} cycloalkyl, group or a linear or branched (C_3-C_{10}) cycloalkyl - (C_1-C_5) alkyl group, or C_6-C_{14} aryl, or a linear or branched (C_6-C_{14}) aryl - (C_1-C_5) alkyl group, or a heterocyclic or linear or branched heterocyclo - (C_1-C_5) alkyl group, said heterocyclic group containing at least a heteroatom selected from the group consisting of nitrogen atom, optionally substituted with a (C_1-C_5) alkyl group, and/or oxygen and/or sulfur; said alkyl, alkenyl, cycloalkyl, cycloalkylalkyl, aryl, aryl-alkyl, heterocyclic or heterocyclo-alkyl groups, being optionally substituted with other groups selected from the group consisting of: halogen, hydroxy, C_1-C_5 alkyl, C_1-C_5 alkoxy, phenyl, cyano, nitro, $-NR_{12}R_{13}$, wherein R_{12} and R_{13}, which may be the same or different, are hydrogen, linear or branched (C_1-C_5) alkyl; a pharmaceutically acceptable ester of the –COOH group; or the–$CONR_{14}R_{15}$ group, wherein R_{14} and R_{15}, which may be the same or different, are hydrogen or linear or branched (C_1-C_5) alkyl; or

R_{10} is a (C_6-C_{10}) aroyl residue optionally substituted by one or more groups selected from the group consisting of: halogen, hydroxy, linear or branched (C_1-C_5) alkyl, C_1-C_5 alkoxy, phenyl, cyano, nitro, $-NR_{16}R_{17}$, wherein R_{16} and R_{17}, which may be the same or different, are hydrogen, linear or branched (C_1-C_8) alkyl;

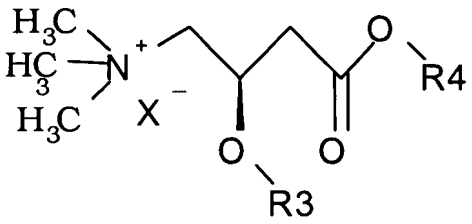
n is the number 0 or 1;

R_{11} is hydrogen, linear or branched C_1-C_5 alkyl, linear or branched C_2-C_5 alkenyl, C_3-C_{10} cycloalkyl, (C_3-C_{10}) cycloalkyl - linear or branched (C_1-C_5) alkyl, C_6-C_{14} aryl, (C_6-C_{14}) aryl - linear or branched alkyl (C_1-C_5);

R_8 and R_9, which may be the same or different are hydrogen, hydroxy, linear or branched C_1-C_5 alkoxy;

their N_1-oxides, their single isomers, in particular the syn and anti isomers of the–$C(R_{11})=N-O_{(n)}R_{10}$ group, their possible enantiomers, diastereoisomers and relative admixtures, the pharmaceutically acceptable salts thereof;

said method comprising encapsulating said antitumor drug into a liposome comprising a compound of formula (II)



(II)

where:

R_3 is a saturated linear or branched acyl chain, with 4-26 carbon atoms;

R_4 is a saturated or unsaturated, linear or branched alkyl chain, with 4-26 carbon atoms; and

X^- is the anion of a pharmacologically acceptable acid, to obtain a liposome containing said antitumor drug, and

administering said liposome to said subject.

95. (Previously Presented) The method according to claim 94, in which R_3 is selected from the group consisting of nonanoyl, dodecanoyl, myristoyl, palmitoyl, stearoyl and oleolyl.

96. (Previously Presented) The method according to claim 94, in which R_4 is selected from the group consisting of nonyl, undecyl, tetradecyl, hexadecyl and oleyl.

97. (Previously Presented) The method according to claim 94, in which X^- is selected from the group consisting of chloride; bromide; iodide; aspartate; acid aspartate; citrate;

acid citrate; tartrate; acid tartrate; phosphate; acid phosphate; fumarate; acid fumarate; glycerophosphate; glucose phosphate; lactate; maleate; acid maleate; mucate; orotate; oxalate; acid oxalate; sulphate; acid sulphate; trichloroacetate; trifluoroacetate; methane sulphonate; pamoate and acid pamoate.

98. (Previously Presented) The method according to claim 94, in which the compound of formula (II) is selected from the group consisting of palmitoyl L-carnitine chloride undecyl ester; stearoyl L-carnitine chloride undecyl ester; stearoyl L-carnitine chloride tetradecyl ester; palmitoyl L-carnitine chloride tetradecyl ester; myristoyl L-carnitine chloride tetradecyl ester; palmitoyl L-carnitine bromide hexadecyl ester, and oleolyl L-carnitine chloride oleyl ester.

99. (Previously Presented) The method according to claim 94, in which said derivative of camptothecin is selected from the group consisting of 7-benzyloxyiminomethylcamptothecin or 7-butoxyiminomethylcam-ptothecin.

100. (Previously Presented) The method according to claim 94, in which the liposome additionally contains helper lipids.

101. (Previously Presented) The method according to claim 100, in which said helper lipid is selected from the group consisting of cholesterol, 1-palmitoyl-2-oleoyl phosphatidyl choline or dioleyl phosphatidyl choline.

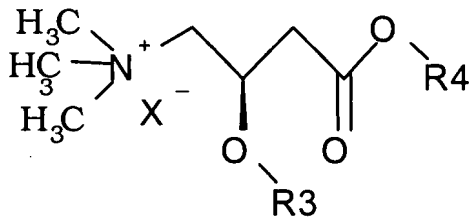
102. (Previously Presented) The method according to claim 94, wherein said antitumor drug is 7-butoxyiminomethylcamptothecin and said liposome comprises the compound palmitoyl L-carnitine undecyl ester.

103. (Previously Presented) The method according to claim 94, wherein said antitumor drug is taxol and said liposome comprises the compound palmitoyl L-carnitine undecyl ester.

104. (Previously Presented) The method according to claim 94, wherein said liposome is administered orally, parenterally, intravenously, intramuscularly, subcutaneously, transdermally or in the form of a nasal or mouth spray.

105. (Previously Presented) The method according to claim 94, wherein lungs are said target organ.

106. (New) A method of topically applying a cosmetic using a liposome comprising a compound of formula (II)



(II)

where:

R_3 is a saturated linear or branched acyl chain, with 4-26 carbon atoms;

R_4 is a saturated or unsaturated, linear or branched alkyl chain, with 4-26 carbon atoms; and

X^- is the anion of a pharmacologically acceptable acid.